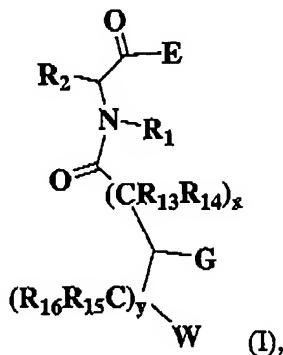


Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application.

Listing of claims:

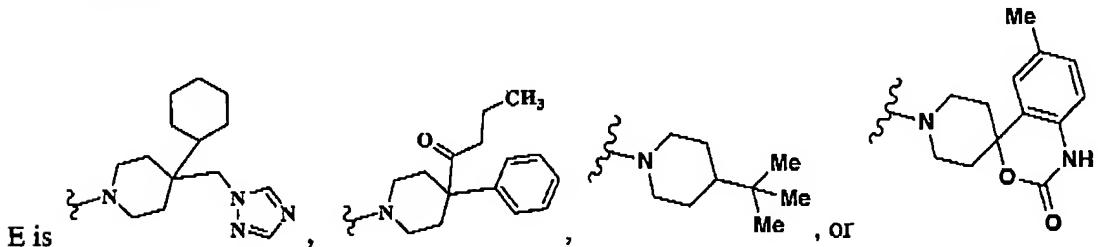
1. (Currently amended) A compound of formula (I),



or a pharmaceutically-acceptable salt or hydrate, thereof, in which:

~~R₁ is hydrogen or C₁₋₆alkyl or is taken together with R₂ or R₃ to form a monocyclic or bicyclic aryl, cycloalkyl, heteroaryl or heterocycle;~~

R₂ is C₁₋₆alkyl or C₂₋₆alkenyl optionally substituted with one to three aryl, cycloalkyl, or heteroaryl, provided that where G is C₂₋₆alkenyl, A₄-NR₁₈CO₂R₁₉, or A₄-SO₂R₁₇, or when y is 0, R₂ ~~is~~ may be C₁₋₆alkyl or C₂₋₆alkenyl, each optionally substituted with heteroaryl;



G is selected from A₄-NR₁₈C(=O)R₁₉, A₄-NR₁₈SO₂R₁₇, A₄-NR₁₈CO₂R₁₉, and

A₄-NR₂₀C(=O)NR₁₈R₁₉ wherein A₄ is a bond, C₁₋₆alkylene, or C₂₋₆alkenylene, or where G is A₄-NR₁₈CO₂R₁₉, or when y is 0, R₂ ~~is~~ may be C₁₋₆alkyl or C₂₋₆alkenyl, each optionally substituted with heteroaryl;

W is selected from substituted or unsubstituted heterocyclo, heteroaryl, or cycloalkyl selected from azetidinyl and imidazolyl, each optionally substituted with lower alkyl;

R_{13} , R_{14} , R_{15} and R_{16} are hydrogen-selected independently of each other from hydrogen, alkyl, substituted alkyl, amino, alkylamine, hydroxy, alkoxy, aryl, cycloalkyl, heteroaryl, or heterocyclo, or R_{13} and R_{14} , or R_{15} and R_{16} , when attached to the same carbon atom, may join to form a spirocycloalkyl ring;

R_{17} is alkyl, substituted alkyl, cycloalkyl, aryl, heterocyclo, or heteroaryl;

R_{18} , R_{19} , and R_{20} are independently selected from hydrogen, alkyl, substituted alkyl, alkenyl, substituted alkenyl, aryl, heteroaryl, cycloalkyl, heterocyclo, or $C(=O)R_{28}$; or when G is $NH(C=O)R_{19}$, R_{19} may be a bond joined to W to define a heterocyclo ring; provided, however, that when y is at least one, W is imidazolyl, indolyl, $-NR_{24}R_{22}$, or $-OR_{23}$, and G is $-NR_{18}C(=O)R_{19}$, then R_{19} is not a C_1 -alkyl having the substituent $-NR_{29}R_{31}$;

R_{29} and R_{31} are selected from hydrogen, alkyl, haloalkyl, hydroxyalkyl, phenylalkyl, and alkoxycarbonylalkyl, or R_{29} and R_{31} taken together form a heterocyclo ring;

x is 0, 1, or 2; and

y is 0, 1, 2, 3 or 4.

2. (Currently amended) A compound according to claim 1, or a pharmaceutically-acceptable salt or hydrate, thereof, in which:

G is selected from:

- a) $-NR_{18}C(=O)R_{19}$;
- b) C_{1-6} alkylene or C_{2-6} alkenylene joined to one of $NR_{18}C(=O)R_{19}$, $NR_{18}CO_2R_{19}$, $NR_{18}SO_2R_{17}$, and $NR_{20}C(=O)NR_{48}R_{49}$;

R_{17} is C_{1-4} alkyl, C_{5-6} cycloalkyl, phenyl, or benzyl;

R_{18} , R_{19} , and R_{20} are independently selected from hydrogen, C_{1-4} alkyl, phenyl, benzyl, C_{5-6} cycloalkyl, $-C(=O)CH_2$ (phenyloxy), $-C(=O)CH_2$ (benzyloxy), imidazolyl, pyridyl, furyl, thiienyl, or C_{1-4} alkyl or C_{2-4} alkenyl substituted with one of phenyl, pyridyl, furyl, cyclopentyl, cyclohexyl, CO_2Me , phenyloxy, or benzyloxy, wherein each ringed group of R_{18} , R_{19} , and R_{20} in turn is optionally substituted with one to two R_{36} , and/or optionally has a benzene ring or five membered heterocyclo having two oxygen atoms fused thereto; and

R₃₆ is halogen, methoxy, nitro, phenyl, phenoxy, or alkylamino.

3. (Currently amended) A compound according to claim 2, or a pharmaceutically-acceptable salt or hydrate, thereof, in which

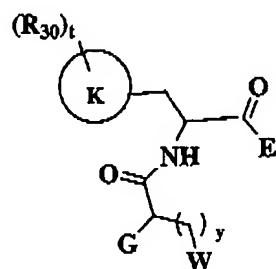
G is $-\text{NR}_{18}\text{C}(=\text{O})\text{R}_{19}$,

R₁₈ is hydrogen or lower alkyl, and

R₁₉ is C₁₋₄alkyl, C₂₋₄alkenyl, phenyl, benzyl, C₅₋₆cycloalkyl, $-\text{C}(=\text{O})\text{CH}_2(\text{phenoxy})$, $-\text{C}(=\text{O})\text{CH}_2(\text{benzyloxy})$, imidazolyl, pyridyl, furyl, thienyl, or C₁₋₄alkyl or C₂₋₄alkenyl substituted with one of phenyl, phenyl, pyridyl, furyl, cyclopentyl, cyclohexyl, CO₂Me, phenoxy, and benzyloxy, wherein each ringed group of R₁₉ in turn is optionally substituted with one to two R₃₆, and/or optionally has a benzene ring or five membered heterocyclo having two oxygen atoms fused thereto.

4. (Previously Presented) A compound according to claim 2, or a pharmaceutically-acceptable salt or hydrate, thereof, in which W is azetidinyl or imidazolyl.

5. (Previously Presented) A compound according to claim 1, or a pharmaceutically-acceptable salt or hydrate, thereof, having the formula:



in which

K is phenyl or thiazolyl;

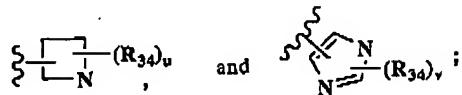
R₃₀ is selected from C₁₋₄alkyl, hydroxy, alkoxy, halogen, nitro, cyano, amino, alkylamino, phenyl, and $-\text{C}(=\text{O})\text{phenyl}$;

t is 0, 1 or 2; and

y is 0, 1 or 2.

6. (Canceled)

7. (Previously presented) A compound according to claim 1, or a pharmaceutically-acceptable salt or hydrate, thereof, in which
W is a ring selected from:



R_{34} at each occurrence is attached to any available carbon or nitrogen atom of W and is selected from
 C_{1-6} alkyl

u is selected from 0, 1, 2, and 3; and

v is 0, 1 or 2.

8. – 9. (Canceled)

10. (Previously presented) A compound according to claim 1, or a pharmaceutically-acceptable salt or hydrate, thereof, in which

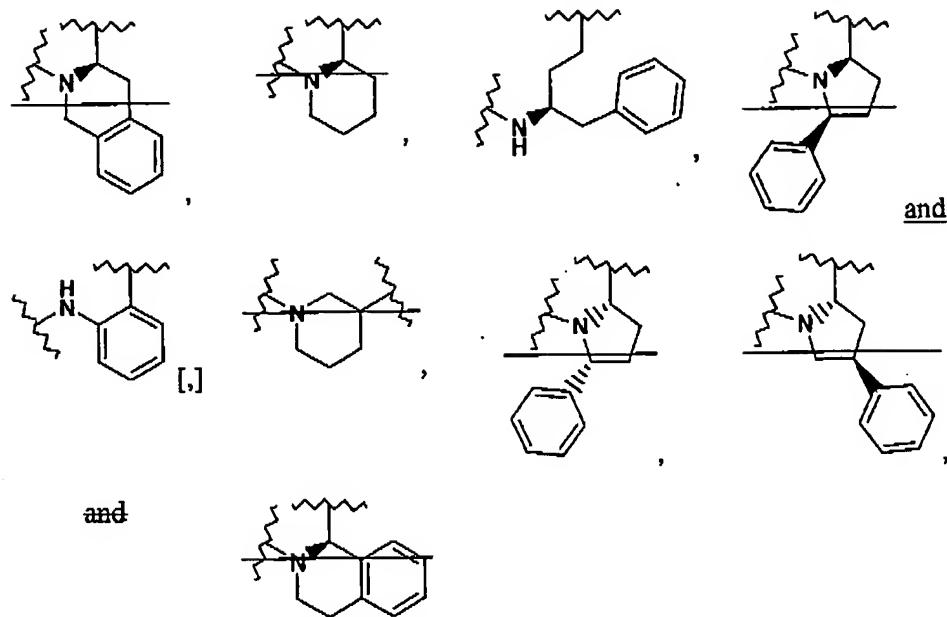
R_2 is selected from C_{1-6} alkyl, C_{2-6} alkenyl, C_{2-6} alkenylene-K, and $-(CH_2)_g-K$;

K is selected from phenyl, naphyl, thienyl, thiazolyl, pyridinyl, pyrimidinyl, and C_{5-6} cycloalkyl,
wherein each group K in turn is optionally substituted with one to three R_{30} or has a benzene
ring fused thereto, which also may be substituted with one to three R_{30} ;

R_{30} is selected from C_{1-4} alkyl, hydroxy, alkoxy, halogen, nitro, cyano, amino, alkylamino, phenyl,
and acylphenyl; and

g is 0, 1, 2 or 3.

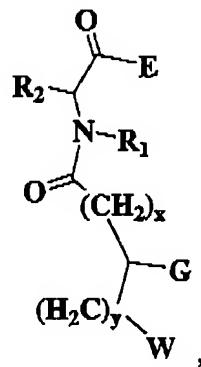
11. (Currently Amended) A compound according to claim 1, or a pharmaceutically-acceptable salt or hydrate, thereof, in which $-N(R_1)-CH(R_2)-$ taken together are selected from,



12. (Previously presented) A compound according to claim 1, or a pharmaceutically-acceptable salt or hydrate, thereof, in which
 R_1 is hydrogen or C_{1-4} alkyl.

13. (Canceled)

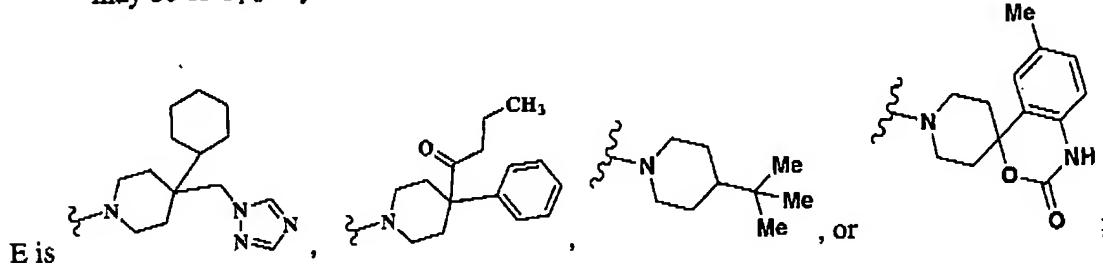
14. (Currently amended) A compound having the formula,



or a pharmaceutically-acceptable salt or hydrate, thereof, in which:

R₁ is hydrogen or C₁₋₆alkyl or is taken together with R₂ or R₃ to form a monocyclic or bicyclic aryl, cycloalkyl, heteroaryl or heterocycle;

R₂ is C₁₋₆alkyl or C₂₋₆alkenyl optionally substituted with one to three aryl, cycloalkyl, or heteroaryl, provided that where G is C₂₋₆alkenyl, or [A₁]-NR₁₈CO₂R₁₉, or A₄-SO₂R₁₇, or when y is 0, R₂ may be or C₁₋₆alkyl or C₂₋₆alkenyl, each optionally substituted with heteroaryl;



G is selected from:

a) NR₁₈C(=O)R₁₉;

b) C₁₋₆alkylene or C₂₋₆alkenylene joined to one of NR₁₈C(=O)R₁₉, NR₁₈CO₂R₁₉, NR₁₈SO₂R₁₇, and NR₂₀C(=O)NR₁₈R₁₉;

W is selected from -substituted or unsubstituted heterocyclo, heteroaryl, or cycloalkyl selected from azetidinyl and imidazolyl, each optionally substituted with lower alkyl;

R₁₇ is alkyl, substituted alkyl, cycloalkyl, aryl, heterocyclo, or heteroaryl;

R₁₈, R₁₉, and R₂₀ are independently selected from hydrogen, alkyl, alkenyl, aryl, heteroaryl, cycloalkyl, heterocyclo, C(=O)R₂₈ or a C₁₋₄alkyl or C₂₋₄alkenyl substituted with one or more of aryl, heteroaryl, cycloalkyl, heterocyclo, alkoxy carbonyl, phenoxy, and/or benzyloxy, and each of said ringed groups of R₁₈, R₁₉, and R₂₀ in turn is optionally substituted with one to two R₃₆;

R₂₁ and R₂₂ are selected from alkyl and substituted alkyl;

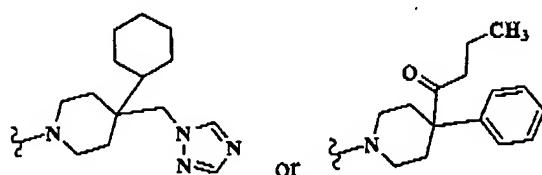
R₃₆ is halogen, methoxy, nitro, phenyl, phenoxy, or alkylamino;

x is 0, 1, or 2; and

y is 0, 1, 2, 3 or 4.

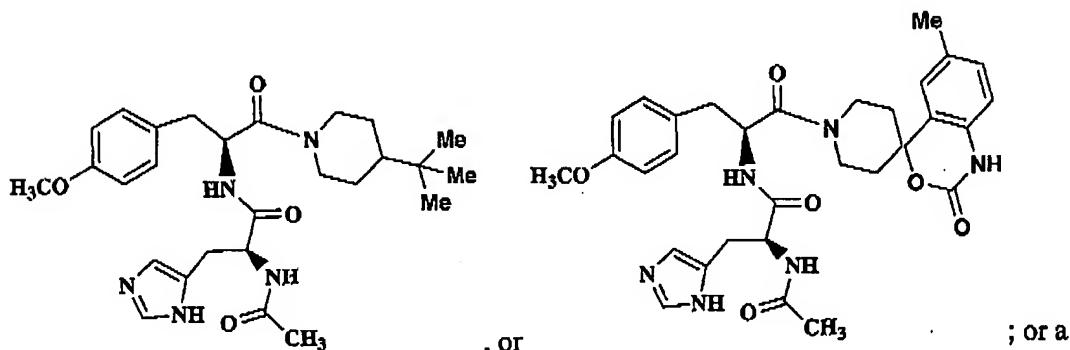
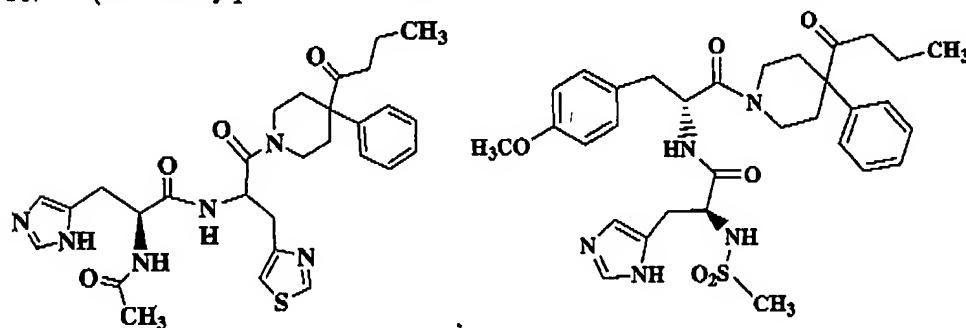
15. (Cancelled)

16. (Previously presented) A compound according to claim 14, or a pharmaceutically-acceptable salt or hydrate, thereof, in which E is



17. (Previously presented) A compound according to claim 14, or a pharmaceutically-acceptable salt or hydrate, thereof, in which G is $\text{NHC}(=\text{O})(\text{alkyl})$ or $\text{NHC}(=\text{O})\text{phenyl}$.

18. (Previously presented) A compound according to claim 1, having the formula,



, pharmaceutically-acceptable salt or hydrate, thereof.

19. (Previously presented) A pharmaceutical composition comprising at least one compound according to claim 1 or a pharmaceutically-acceptable salt or hydrate, thereof; and a pharmaceutically-acceptable carrier or diluent.

20. - 23. (Canceled)